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- 1. An adjuvant comprising a surfactant and the apolar fraction or part of the apolar fraction of the total lipid extract of a mycobacterium, e.g. the BCG, *M.microti, M.tuberculosis* and *M.vaccae*.
- An adjuvant according to claim 1 where the part of the apolar fraction of the lipid extract can be phthiocerol dimycocerosates, trehalose mycolipenates, glycosylated phenol phthiocerols (including phenolic glycolipids, PGL's), trehalose mycolates, sulfolipids, triacylglycerols or menaquinones
- 3. An adjuvant according to claim 1-2 where the surfactant is cationic.
- An adjuvant according to claim 3 where the surfactant is DDA, DODA, DC-chol or DOTAP.
- An adjuvant according to claim 1-2 where the surfactant is neutral or anionic, e.g. DOPE/PC or DOPE/PC/PG.
- 6. A vaccine comprising an adjuvant according to claim 1-5.
- A vaccine according to claim 6 for parenterally, oral or mucosal administration.
- 8. A vaccine according to claim 7 where the antigenic component comprises an antigenic epitope from a virulent mycobacterium, e.g. *Mycobacterium tuberculosis*, *M. bovis* or *M.africanum*.
- A vaccine according to claim 8 where the antigenic component is an ESAT6-Ag85B hybrid or a fragment hereof.
- A vaccine according to claim 7 for treating cancer, allergy or autoimmune diseases.
- 11. A delivery system comprising an adjuvant according to claim 1-5.

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12. Preparing an adjuvant according to claim 1-5 using thin lipid film method.